FILE 'HOME' ENTERED AT 15:10:32 ON 24 OCT 2006 => file biosis medline caplus wpids uspatfull COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 0.21 0.21 FILE 'BIOSIS' ENTERED AT 15:10:53 ON 24 OCT 2006 Copyright (c) 2006 The Thomson Corporation FILE 'MEDLINE' ENTERED AT 15:10:53 ON 24 OCT 2006 FILE 'CAPLUS' ENTERED AT 15:10:53 ON 24 OCT 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS) FILE 'WPIDS' ENTERED AT 15:10:53 ON 24 OCT 2006 COPYRIGHT (C) 2006 THE THOMSON CORPORATION FILE 'USPATFULL' ENTERED AT 15:10:53 ON 24 OCT 2006 CA INDEXING COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS) *** YOU HAVE NEW MAIL *** => s polycationic (3a) multichromophore? 11 POLYCATIONIC (3A) MULTICHROMOPHORE? => s ll and peptide nucleic acid? 8 L1 AND PEPTIDE NUCLEIC ACID? L2=> dup rem 12 PROCESSING COMPLETED FOR L2 L38 DUP REM L2 (0 DUPLICATES REMOVED) => d 13 bib abs 1-8 L3 ANSWER 1 OF 8 CAPLUS COPYRIGHT 2006 ACS on STN 2006:681478 CAPLUS AN 145:138599 DN Cationic conjugated polymers suitable for strand-specific polynucleotide TI detection in homogeneous and solid state assays Bazan, Guillermo C.; Liu, Bin IN PA The Regents of the University of California, USA PCT Int. Appl., 71 pp. SO CODEN: PIXXD2 DΤ Patent LA English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE -----____ ----------PΤ WO 2006074471 20060713 WO 2006-US882 A2 20060110 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE,

SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC,

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,

VN, YU, ZA, ZM, ZW

KG, KZ, MD, RU, TJ, TM

US 2006183140 20060817 **A1** US 2006-329495 20060110 PRAI US 2005-642901P P 20050110 The invention further relates to polycationic multichromophores, which may be conjugated polymers, and methods, articles and compns. employing them as described herein. In some aspects, the invention relates to methods, articles and compns. for the detection and anal. of biomols. in a sample. Provided assays include those determining the presence of a target biomol. in a sample or its relative amount, or the assays may be quant. or semi-quant. The methods can be performed on a substrate. The methods can be performed in an array format on a substrate, which can be a sensor. In some embodiments, detection assays are provided employing sensor biomols. that do not comprise a fluorophore that can exchange energy with the cationic multichromophore. In some aspects biol. assays are provided in which energy is transferred between one or more of the multichromophore, a label on the target biomol., a label on the sensor biomol., and/or a fluorescent dye specific for a polynucleotide, in all permutations. The multichromophore may interact at least in part electrostatically with the sensor and/or the target, and an increase in energy transfer with the polymer may occur upon binding of the sensor and the target. Other variations of the inventions are described further herein. Thus, poly[9,9'-bis((6''-N,N,Ntrimethylammonium)hexyl)fluorene-co-alt-4,7-(2,1,3benzothiadiazole)dibromide] (PFBT) in phosphate buffer containing 5% 1-methyl-2-pyrrolidinone was combined with Cy5-labeled PNA and its complementary target DNA. Excitation at 460 nm resulted in intense red emission from Cy5. There was no energy transfer for the solution containing the Cy5-labeled PNA and target DNA. To prepare PFBT, 2,7-bis[9,9'-bis(6''bromohexyl)fluorenyl]-4,4,5,5-tetramethyl-[1,3,2]dioxaborolane was first synthesized from 9,9'-bis(bromohexyl)-2,7-dibromofluorene and 2-isopropoxy-4,4,5,5-tetramethyl-[1.3.2]-dixoaborolane. Suzuki copolymn. of 2,7-bis[9,9'-bis(6''-bromohexyl)fluorenyl]-4,4,5,5-tetramethyl-[1,3,2]dioxaborolane and 4,7-dibromo-2,1,3-benzothiazole produced the PFBT precursor. The polymer PFBT was prepared from the precursor polymer, poly[9,9'-bis((6'-bromohexyl)fluorene)-co-alt-4,7-(2,1,3benzothiadiazole)], by reaction with Me3N. L3 ANSWER 2 OF 8 USPATFULL on STN 2006:254283 USPATFULL ΑN ΤI Methods and articles for strand-specific polynucleotide detection with cationic multichromophores TN Bazan, Guillermo C., Santa Barbara, CA, UNITED STATES Liu, Bin, Singapore, SINGAPORE PA The Regents of the University of California, Oakland, CA, UNITED STATES (U.S. corporation) US 2006216734 PΙ **A**1 20060928 AΙ US 2006-329861 **A**1 20060110 (11) US 2005-642883P 20050110 (60) PRAI DTUtility APPLICATION FS FITCH EVEN TABIN AND FLANNERY, 120 SOUTH LA SALLE STREET, SUITE 1600, CHICAGO, IL, 60603-3406, US CLMN Number of Claims: 48 ECL Exemplary Claim: 1 12 Drawing Page(s) LN.CNT 2161 CAS INDEXING IS AVAILABLE FOR THIS PATENT. The invention further relates to polycationic multichromophores, which may be conjugated polymers, and methods, articles and compositions employing them as described herein. In some aspects, the invention relates to methods, articles and compositions for the detection and analysis of biomolecules in a sample.

Provided assays include those determining the presence of a target biomolecule in a sample or its relative amount, or the assays may be quantitative or semi-quantitative. The methods can be performed on a substrate. The methods can be performed in an array format on a substrate, which can be a sensor. In some embodiments, detection assays are provided employing sensor biomolecules that do not comprise a fluorophore that can exchange energy with the cationic multichromophore. In some aspects biological assays are provided in which energy is transferred between one or more of the multichromophore, a label on the target biomolecule, a label on the sensor biomolecule, and/or a fluorescent dye specific for a polynucleotide, in all permutations. The multichromophore may interact at least in part electrostatically with the sensor and/or the target, and an increase in energy transfer with the polymer may occur upon binding of the sensor and the target. Other variations of the inventions are described further herein.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L3
     ANSWER 3 OF 8 USPATFULL on STN
AN
       2006:240539 USPATFULL
ΤI
      Methods and compositions for aggregant detection
TN
      Bazan, Guillermo C., Santa Barbara, CA, UNITED STATES
      Liu, Bin, Singapore, SINGAPORE
PA
      The Regents of University of California, Oakland, CA, UNITED STATES
       (U.S. corporation)
PΙ
      US 2006204984
                          A1
                               20060914
      US 2006-344942
                               20060131 (11)
ΑI
                          A1
      US 2005-649024P
PRAI
                          20050131 (60)
DТ
      Utility
FS
      APPLICATION
LREP
      FITCH EVEN TABIN AND FLANNERY, 120 SOUTH LA SALLE STREET, SUITE 1600,
      CHICAGO, IL, 60603-3406, US
CLMN
      Number of Claims: 28
ECL
      Exemplary Claim: 1
DRWN
       9 Drawing Page(s)
LN.CNT 2187
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      This invention relates to an aggregation sensor useful for the detection
      and analysis of aggregants in a sample, and methods, articles and
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This invention relates to an aggregation sensor useful for the detection and analysis of aggregants in a sample, and methods, articles and compositions relating to such a sensor. The sensor comprises first and second optically active units, where energy may be transferred from an excited state of the first optically active unit to the second optically active unit. The second optically active unit is present in a lesser amount, but its relative concentration is increased upon aggregation, increasing its absorption of energy from the first optically active units. This increase in energy transfer can be detected in variety of formats to produce an aggregation sensing system for various aggregants, including for quantitation. Other variations of the inventions are described further herein.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L3
     ANSWER 4 OF 8 USPATFULL on STN
       2006:214985
AN
                    USPATFULL
TI
       Cationic conjugated polymers suitable for strand-specific polynucleotide
       detection in homogeneous and solid state assays
IN
       Bazan, Guillermo C., Santa Barbara, CA, UNITED STATES
       Liu, Bin, Singapore, SINGAPORE
{\tt PA}
       The Regents of the University of California, Oakland, CA, UNITED STATES
       (U.S. corporation)
PΙ
       US 2006183140
                          A1
                                20060817
       US 2006-329495
ΑI
                          A1
                                20060110 (11)
       US 2005-642901P
PRAI
                           20050110 (60)
DT
       Utility
FS
       APPLICATION
       FITCH EVEN TABIN AND FLANNERY, 120 SOUTH LA SALLE STREET, SUITE 1600,
LREP
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CHICAGO, IL, 60603-3406, US

CLMN Number of Claims: 63 ECL Exemplary Claim: 1

DRWN 12 Drawing Page(s)

LN.CNT 2181

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention further relates to polycationic multichromophores, which may be conjugated polymers, and methods, articles and compositions employing them as described herein. In some aspects, the invention relates to methods, articles and compositions for the detection and analysis of biomolecules in a sample. Provided assays include those determining the presence of a target biomolecule in a sample or its relative amount, or the assays may be quantitative or semi-quantitative. The methods can be performed on a substrate. The methods can be performed in an array format on a substrate, which can be a sensor. In some embodiments, detection assays are provided employing sensor biomolecules that do not comprise a fluorophore that can exchange energy with the cationic multichromophore. In some aspects biological assays are provided in which energy is transferred between one or more of the multichromophore, a label on the target biomolecule, a label on the sensor biomolecule, and/or a fluorescent dye specific for a polynucleotide, in all permutations. The multichromophore may interact at least in part electrostatically with the sensor and/or the target, and an increase in energy transfer with the polymer may occur upon binding of the sensor and the target. Other variations of the inventions are described further herein.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 5 OF 8 USPATFULL on STN

AN 2005:4283 USPATFULL

TI Methods and compositions for detection and analysis of polynucleotide-binding protein interactions using light harvesting multichromophores

IN Bazan, Guillermo C., Santa Barbara, CA, UNITED STATES
Wang, Shu, Goleta, CA, UNITED STATES
Liu, Bin, Goleta, CA, UNITED STATES

PA The Regents of the University of California (U.S. corporation)

PI US 2005003386 A1 20050106 AI US 2004-779412 A1 20040213 (10)

PRAI US 2003-447860P 20030213 (60)

DT Utility

FS APPLICATION

LREP Bingham McCutchen LLP, Suite 1800, Three Embarcadero Center, San Francisco, CA, 94111-4067

CLMN Number of Claims: 30 ECL Exemplary Claim: 1

DRWN 9 Drawing Page(s)

LN.CNT 1330

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Methods, compositions and articles of manufacture for assaying a sample for a target polynucleotide are provided. A sample suspected of containing the target polynucleotide is contacted with a polycationic multichromophore and a sensor PBP that can bind to the target polynucleotide. The sensor PBP comprises a signaling chromophore to absorb energy from the excited multichromophore and emit light in the presence of the target polynucleotide. The methods can be used in multiplex form. Kits comprising reagents for performing such methods are also provided.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L3 ANSWER 6 OF 8 USPATFULL on STN

AN 2004:280256 USPATFULL

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Methods and compositions for detection and analysis of polynucleotides
       using light harvesting multichromophores
IN
       Bazan, Guillermo C., Santa Barbara, CA, UNITED STATES
       Gaylord, Brent S., Santa Barbara, CA, UNITED STATES
PA
       The Regents of the University of California, Oakland, CA, UNITED STATES
       (U.S. corporation)
PΙ
       US 2004219556
                          A1
                               20041104
ΑI
       US 2003-600286
                          A1
                               20030620 (10)
PRAI
       US 2002-406266P
                           20020826 (60)
DT
       Utility
FS
       APPLICATION
       David W. Maher, Bingham McCutchen LLP, 28th Floor, Three Embarcadero
LREP
       Center, San Francisco, CA, 94111
       Number of Claims: 30
CLMN
ECL
       Exemplary Claim: 1
DRWN
       4 Drawing Page(s)
LN.CNT 1178
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Methods, compositions and articles of manufacture for assaying a sample
       for a target polynucleotide are provided. A sample suspected of
       containing the target polynucleotide is contacted with a
       polycationic multichromophore and a sensor PNA
       complementary to the target polynucleotide. The sensor PNA comprises a
       signaling chromophore to absorb energy from the excited multichromophore
       and emit light in the presence of the target polynucleotide. The methods
       can be used in multiplex form. Kits comprising reagents for performing
       such methods are also provided.
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 7 OF 8 USPATFULL on STN
T.3
       2004:184461 USPATFULL
AN
TТ
       Methods and compositions for detection and analysis of polynucleotides
       using light harvesting multichromophores
IN
       Bazan, Guillermo C., Santa Barbara, CA, UNITED STATES
       Gaylord, Brent S., Santa Barbara, CA, UNITED STATES
       Wang, Shu, Goleta, CA, UNITED STATES
PA
       The Regents of the University of California, Oakland, CA, UNITED STATES
       (U.S. corporation)
       US 2004142344
PΙ
                          Α1
                               20040722
       US 2003-648945
ΑI
                          A1
                               20030826 (10)
       US 2002-406266P
PRAI
                          20020826 (60)
DT
       Utility
       APPLICATION
FS
LREP
       BINGHAM, MCCUTCHEN LLP, THREE EMBARCADERO, SUITE 1800, SAN FRANCISCO,
       CA, 94111-4067
CLMN
       Number of Claims: 55
ECL
       Exemplary Claim: 1
DRWN
       5 Drawing Page(s)
LN.CNT 1305
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
       Methods, compositions and articles of manufacture for assaying a sample
       for a target polynucleotide are provided. A sample suspected of
       containing the target polynucleotide is contacted with a
       polycationic multichromophore and a sensor
       polynucleotide complementary to the target polynucleotide. The sensor
       polynucleotide comprises a signaling chromophore to receive energy from
       the excited multichromophore and increase emission in the presence of
       the target polynucleotide. The methods can be used in multiplex form.
       Kits comprising reagents for performing such methods are also provided.
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.

TΙ

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AN
     2004-142830 [14]
                        WPIDS
CR
     2004-784599
DNC
    C2004-057433 [14]
DNN
    N2004-113890 [14]
    Assay method, by contacting sample e.g. blood, urine with sensor
ΤI
    peptide nucleic acid which has signaling
    chromophore to absorb energy from excited multichromophore and emit light
     in presence of target polynucleotide
DC
    A89; B04; D16; S03
IN
    BAZAN G C; GAYLORD B S; WANG S
PA
     (REGC-C) UNIV CALIFORNIA
CYC
    103
PIA WO 2004001379
                    A2 20031231 (200414)* EN
                                               34[4]
    AU 2003243722
                    A1 20040106 (200447)
    US 20040142344 A1 20040722 (200449)
    US 20040219556 A1 20041104 (200473)
    KR 2005010956 A 20050128 (200535)
                                          KO
    EP 1534857
                    A2 20050601 (200536)
                                          EN
    JP 2005530182
                    W 20051006 (200566)
                                          JA
                                               28
    CN 1675377
                    A 20050928 (200610)
                                          zH
    ZA 2005000529
                    A 20051228 (200612) EN
                                              43
    CN 1694967
                    A 20051109 (200618) ZH
ADT WO 2004001379 A2 WO 2003-US19678 20030620; US 20040142344 A1 Provisional
    US 2002-406266P 20020826; US 20040219556 A1 Provisional US 2002-406266P
    20020826; AU 2003243722 A1 AU 2003-243722 20030620; CN 1675377 A CN
    2003-819836 20030620; EP 1534857 A2 EP 2003-761235 20030620; US
    20040219556 A1 US 2003-600286 20030620; EP 1534857 A2 WO 2003-US19678
    20030620; JP 2005530182 W WO 2003-US19678 20030620; US 20040142344 A1 US
    2003-648945 20030826; JP 2005530182 W JP 2004-516105 20030620; KR
    2005010956 A KR 2004-720729 20041220; ZA 2005000529 A ZA 2005-529
    20050119; CN 1694967 A CN 2003-824651 20030826
FDT AU 2003243722 Al Based on WO 2004001379 A; EP 1534857 A2 Based on WO
    2004001379 A; JP 2005530182 W Based on WO 2004001379 A
PRAI US 2002-406266P 20020826
    US 2002-390524P 20020620
    US 2003-600286 20030620
    US 2003-648945 20030826
AN
    2004-142830 [14]
                       WPIDS
CR
    2004-784599
AB
    WO 2004001379 A2
                       UPAB: 20060121
     NOVELTY - An assay method, comprising contacting the sample containing
    target polynucleotide with sensor peptide nucleic
    acid (PNA) and polycationic multichromophore
    in a solution under conditions in which sensor PNA can hybridize to target
    polynucleotide.
            DETAILED DESCRIPTION - An assay method, comprising contacting the
    sample containing target polynucleotide with sensor peptide
    nucleic acid (PNA) and polycationic
    multichromophore in a solution under conditions in which sensor
    PNA can hybridize to target polynucleotide, if present, applying a light
    source to solution that can excite multichromophore, and detecting whether
    light is emitted from signaling chromophore of PNA, is new.
           An assay method (M1), comprising:
            (a) providing a sample that is suspected of containing a target
    polynucleotide;
            (b) providing a polycationic multichromophore
    that electrostatically interacts with the target polynucleotide and upon
    excitation is capable of transferring energy to a signaling chromophore;
            (c) providing a sensor peptide nucleic
    acid (PNA) (I) that is single-stranded and is complementary to the
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(d) contacting the sample with (I) and the multichromophore in a solution under conditions in which the sensor PNA can hybridize to the

target polynucleotide, the sensor PNA conjugated to the signaling

chromophore;

target polynucleotide, if present;

- (e) applying a light source to the solution that can excite the multichromophore; and
- (f) detecting whether light is emitted from the signaling chromophore.

INDEPENDENT CLAIMS are also included for the following:

- (1) a polynucleotide sensing solution comprising (I), a polycationic multichromophore (II) that can electrostatically interact with the phosphate backbone of the target polynucleotide and is capable of transferring energy to the signaling chromophore upon excitation when brought into proximity to it upon hybridization of the sensor PNA to the target polynucleotide; and
- (2) a kit for assaying a sample for a target polynucleotide comprising (I) and (II).

USE - (M1) is useful for assaying target polynucleotide in a sample. The target polynucleotide is DNA or RNA. The sample is comprises single, or double stranded target polynucleotide. The target polynucleotide is produced by an amplification reaction. (All claimed.) (M1) is useful for assaying target nucleic acid in sample such as blood, urine, milk, semen, sputum, mucus, buccal swab, vaginal swab, rectal swab, aspirate, needle biopsy, etc.

ADVANTAGE - (M1) allows analysis of target polynucleotide that occurs naturally in the sample or can be amplified prior to or in conjugation with analysis. By using multiple different sensor PNAs, multiple different polynucleotides can be independently detected and assayed.

DESCRIPTION OF DRAWINGS - The drawing shows assay method of target polynucleotide in a sample, employing a polycationic polymer as a light harvesting multichromophore.